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CardinalHealth

XIII. SUMMARY OF SAFETY AND EFFECTIVENESS

K052568

510(k) SUMMARY OF SAFETY AND EFFECTIVENESS VINYL POWDER-FREE EXAMINATION GLOVES WITH COATING

Applicant/Sponsor: Cardinal Health
1500 Waukegan Road
McGaw Park, IL 60085

Regulatory Affairs Contact: Amy Hoyd
Cardinal Health
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McGaw Park, IL 60085

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Date Summary Prepared: 03 July 2005

Product Trade Name: Esteem Stretchy Synthetic with Neu-Thera

Common Name: Examination Glove

Classification: Patient Examination Glove

Predicate Devices: Vinyl Powder-Free Examination Gloves, Grand Work Plastic Products Co., Ltd.

Description: Vinyl Powder-Free Examination Gloves with coating are formulated using Vinyl and offered powder-free.

Intended Use: These examination gloves are disposable devices intended for medical purposes that are worn on the examiner's hands or fingers to prevent contamination between patient and examiner.



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Substantial Equivalence: Esteem Stretchy Synthetic with Neu-Thera are substantially equivalent to Grand Work Plastic Products Co., Ltd's Vinyl Powder-Free Examination Gloves in that they provide the following characteristics:

- same intended use
- same sizes
- both made of Vinyl
- both offered beaded and powder-free

Summary of Testing:

<u>Test</u>	<u>Result</u>
Primary Skin Irritation	Gloves show no irritation.
Guinea Pig Maximization	Gloves do not display any potential for irritation.
Tensile Strength	Gloves meet or exceed requirements per ASTM D5250-00e4.
Barrier Defects	Gloves meet or exceed requirements per 21 CFR§800.20 and ASTM D5250-00e4.

**DEPARTMENT OF HEALTH & HUMAN SERVICES**

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV 28 2005

Cardinal Health
C/O Mr. Neil Devine
Responsible Third Party Official
Intertek Testing Services
70 Codman Hill Road
Boxborough, Massachusetts 01719

Re: K052568

Trade/Device Name: Esteem Stretchy Synthetic with Neu-Thera
Regulation Number: 21 CFR 880.6250
Regulation Name: Patient Examination Glove
Regulatory Class: I
Product Code: LYZ
Dated: November 21, 2005
Received: November 22, 2005

Dear Mr. Devine:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

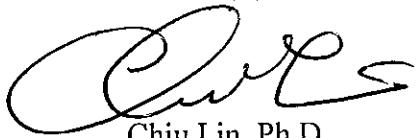
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0115. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Chiu Lin, Ph.D.

Director

Division of Anesthesiology, General Hospital,

Infection Control and Dental Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure



Cardinal Health

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510(k) Number: K052568

Device Name: Esteem Stretchy Synthetic with Neu-Thera

Indications For Use: A patient examination glove is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

Prescription Use _____ AND/OR Over-The Counter Use X _____
(21 CFR 801 Subpart D) (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE)

Concurrence of CDRH, Office of Device Evaluation (ODE)

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